





## Why Solve this Collaboratively?

- The scope of reuse appears very broad both by type of device and facility
- The science base on reuse still not rich but growing we need to hear from experts
- We need to listen to our stakeholders
- Resources
- Public health issues are a joint responsibility

## Science, Compliance, and Surveillance

- Science: FDA needs to base its decisions on science: we need more than we have
- Compliance: Inspections will likely increase, but we will need to partner to work effectively with the hospital sector
- Surveillance: FDA's tools for surveillance of this topic face significant challenges

## Vision for the Future Current Reality Future Vision

- Widespread practice with little data on safety or effectiveness
- Single use labels not clearly meaningful
- Single use labels don't identify vulnerabilities
- Patients are not informed experimentation?

- FDA regulatory approach will be RISK and SCIENCE based
- Single use labels will have clinical relevance
- Single use labels will identify vulnerabilities
- Horizontal and vertical standards critical
- Work with all parties